## In the Claims:

## 1.-2. Cancelled

3. (Currently amended) The A compound of Formula III Claim 1, or a pharmaceutically acceptable salt or stereoisomer thereof, of the Formula III:

$$R^{3a}$$
  $O$   $R^{4a}$   $R^{4b}$   $R^{3b}$   $R^{2a}$   $R^{2a}$   $R^{4b}$ 

wherein:

m is 0, 1 or 2; p' is 0 to 2; r is 0 or 1; s is 0 or 1;

 $R^2$  is (C<sub>1</sub>-C<sub>6</sub>)alkylene-NR<sup>6</sup>R<sup>7</sup>; said alkylene is optionally substituted with up to three substituents selected from OH, (C<sub>1</sub>-C<sub>6</sub>)alkoxy, halogen, CO<sub>2</sub>H, CN, O(C=O)C<sub>1</sub>-C<sub>6</sub> alkyl, oxo, and NR<sup>6</sup>R<sup>7</sup>;

R<sup>2a</sup> is selected from: halogen and (C<sub>1</sub>-C<sub>6</sub>)alkyl;

R<sup>3a</sup> and R<sup>3b</sup> are independently selected from: hydrogen, halogen, and (C<sub>1</sub>-C<sub>6</sub>)alkyl;

 $R^{4a}$  and  $R^{4b}$  are independently selected from: hydrogen, halogen and (C1-C6)alkyl, provided that at lease one is not hydrogen, or

R4a and R4b are combined to form a diradical selected from CH2CH2CH2CH2-,

## -CH2CH2CH2-, -CH-CH-O and -CH-CH-N-;

## R<sup>5</sup> is selected from:

- 1) (C=O)<sub>F</sub>O<sub>S</sub>(C<sub>1</sub>-C<sub>10</sub>)alkyl,
- 2) O<sub>r</sub>(C<sub>1</sub>-C<sub>3</sub>)perfluoroalkyl,
- 3) (C<sub>0</sub>-C<sub>6</sub>)alkylene S(O)<sub>m</sub>R<sup>a</sup>,
- 4)——oxo,
- 5)——OH,
- 6) halo,
- 7) CN,
- 8) (C=O)<sub>F</sub>O<sub>S</sub>(C<sub>2</sub>-C<sub>10</sub>)alkenyl,
- 9) (C=O)<sub>r</sub>O<sub>s</sub>(C<sub>2</sub>-C<sub>10</sub>)alkynyl,
- 10) (C=O)<sub>r</sub>O<sub>S</sub>(C<sub>3</sub>-C<sub>6</sub>)cycloalkyl,
- 11) (C=O)<sub>f</sub>O<sub>S</sub>(C<sub>0</sub>-C<sub>6</sub>)alkylene aryl,
- 12) (C=O)<sub>F</sub>O<sub>S</sub>(C<sub>O</sub>-C<sub>6</sub>)alkylene-heterocyclyl,
- 13) (C=O)rOs(C0-C6)alkylene-N(Rb)2,
- 14)  $C(O)R^a$
- 15) (C<sub>0</sub>-C<sub>6</sub>)alkylene-CO<sub>2</sub>R<sup>a</sup>.
- 16) C(O)H,
- 17) (C<sub>0</sub>-C<sub>6</sub>)alkylene-CO<sub>2</sub>H, and
- 18) C(O)N(Rb)2,

said alkyl, alkenyl, alkynyl, cycloalkyl, aryl, and heterocyclyl is optionally substituted with up to three substituents selected from R<sup>b</sup>, OH, (C<sub>1</sub>-C<sub>6</sub>)alkoxy, halogen, CO<sub>2</sub>H, CN, O(C=O)C<sub>1</sub>-C<sub>6</sub> alkyl, oxo, and N(R<sup>b</sup>)<sub>2</sub>;

 $R^6$  and  $R^7$  are independently selected from:

- 1) H
- 2)  $(C=O)O_bC_1-C_{10}$  alkyl,
- 3) (C=O)ObC3-C8 cycloalkyl,
- 4) (C=O)Obaryl,
- 5) (C=O)Obheterocyclyl,
- 6) C<sub>1</sub>-C<sub>10</sub> alkyl,
- 7) aryl,
- 8) C2-C<sub>10</sub> alkenyl,

- 9)  $C_2$ - $C_{10}$  alkynyl,
- 10) heterocyclyl,
- 11) C3-C8 cycloalkyl,
- 12) SO<sub>2</sub>Ra, and
- 13)  $(C=O)NRb_2$ ,

said alkyl, cycloalkyl, aryl, heterocylyl, alkenyl, and alkynyl is optionally substituted with one or more substituents selected from R<sup>5</sup>, or

R<sup>6</sup> and R<sup>7</sup> can be taken together with the nitrogen to which they are attached to form a monocyclic or bicyclic heterocycle with 4-7 members in each ring and optionally containing, in addition to the nitrogen, one or two additional heteroatoms selected from N, O and S<del>, said monocyclic or bicyclic heterocycle optionally substituted with one or more substituents selected from R<sup>5</sup>;</del>

Ra is (C1-C6)alkyl, (C3-C6)cycloalkyl, aryl, or heterocyclyl; and

Rb is H, (C1-C6)alkyl, (C1-C6)alkyl-NRa2, (C1-C6)alkyl-NH2, (C1-C6)alkyl-NHRa, aryl, heterocyclyl, (C3-C6)cycloalkyl, (C=O)OC1-C6 alkyl, (C=O)C1-C6 alkyl or S(O)2Ra.

4. (Currently amended) The compound according to Claim 3 or a pharmaceutically acceptable salt or stereoisomer thereof, wherein: p', R<sup>2a</sup>, R<sup>3a</sup>, R<sup>3b</sup>, R<sup>4a</sup>, and R<sup>4b</sup> and R<sup>5</sup> are as defined for Formula III and

R<sup>2</sup> is (C<sub>1</sub>-C<sub>6</sub>)alkylene-NR<sup>6</sup>R<sup>7</sup>;

 $R^6$  and  $R^7$  are independently selected from:

- 1) H,
- 2) C<sub>1</sub>-C<sub>10</sub> alkyl,
- 3) aryl,
- 4) heterocyclyl,
- 5) C2-C<sub>10</sub> alkenyl,
- 6) C2-C<sub>10</sub> alkynyl, and
- 7) C3-C8 cycloalkyl,

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said alkyl, cycloalkyl, aryl, heterocylyl, alkenyl, and alkynyl is optionally substituted with one or more substituents selected from R<sup>5</sup>, or

R<sup>6</sup> and R<sup>7</sup> can be taken together with the nitrogen to which they are attached to form a monocyclic or bicyclic heterocycle with 4-7 members in each ring and optionally containing, in addition to the nitrogen, one or two additional heteroatoms selected from N, O and S, said monocyclic or bicyclic heterocycle optionally substituted with one or more substituents selected from R<sup>5</sup>.

5. (Original) A compound which is:

2-(2-bromophenyl)-3-(4-methylphenyl)thieno[2,3-d]pyrimidin-4(3H)-one.

- 6. (Currently amended) A pharmaceutical composition that is comprised of a compound in accordance with Claim + 3 and a pharmaceutically acceptable carrier.
- 7. (Original) A pharmaceutical composition that is comprised of a compound in accordance with Claim 3 and a pharmaceutically acceptable carrier.
  - 8.-11. Cancelled
  - 12.-20. Previously Cancelled
  - 21.-24. Cancelled.
  - 25.-27. Previously Cancelled.
  - 28.-29. Cancelled.
  - 30. Previously Cancelled.
  - 31.-34. Cancelled.